

AMENDMENT

Amendments to the Claims:

Please amend the claims as follows, without prejudice:

In the Claims:

1-45 (Canceled).

46. (Currently Amended) A serum-stable amphoteric liposomal formulation[[s]] comprising a liposome with an aqueous interior and at least one active substance in the aqueous interior, characterized in

that wherein the liposomes comprise

- neutral lipids with a membrane proportion of 10 to 60 mole-%,
- cholesterol with a proportion of 30 to 50 mole-%,

and, as charged lipids, either

- amphoteric lipids with a proportion of 5 to 30 mole-%,
- or
- mixtures of cationic and anionic lipids with an overall proportion of 50 mole% at maximum,

and that wherein the active substance comprises at least one oligonucleotide.

47. (Currently Amended) The liposomal formulation according to claim 46, characterized in that wherein the proportion of cholesterol is 35 to 45 mole-%[[,]] and the proportion of amphoteric lipids is 5 to 20 mole-% and/or the proportion of said mixtures is 15 to 45 mole-%.

48. (Currently Amended) The liposomal formulation according to claim 46, characterized in that wherein the oligonucleotides are constituted of 5-100, preferably 5-40 and more preferably 10-25 deoxyribonucleotides, ribonucleotides or chemically modified derivatives thereof.

49. (Currently Amended) The liposomal formulation according to claim 146, characterized in that wherein the oligonucleotides are present as single strands, double strands, or in complex folding.

50. (Currently Amended) The liposomal formulation according to claim 46, characterized in that wherein the single strands are present as oligonucleotide is an antisense oligonucleotides, the double strands as small interfering RNA and/or decoy oligonucleotides and/or the complex foldings as aptamers and/or spiegelmers.

51. (Currently Amended) The liposomal formulation according to claim 146, characterized in that wherein the oligonucleotide is an aptamer.

52. (Currently Amended) The liposomal formulation according to claim 146, characterized in that wherein the oligonucleotide is a spiegelmer.

53. (Currently Amended) The liposomal formulation according to claim 146, characterized in that wherein the liposomal membrane liposome has the a molar composition selected from the group consisting of:

DMPC/MoChol/DMPS/Chol 40:10:10:40,  
DMPC/AC/Chol 50:10:40,  
DMPC/HisChol/DPPS/Chol 35:10:15:40,  
DMPC/IsohistsuccDG/Chol 50:10:40,  
DMPC/MoChol/DGSucc/Chol 35:10:15:40,  
DMPC/MoChol/DGSucc/Chol 40:10:10:40,  
POPC/MoChol/DGSucc/Chol 35:10:15:40,  
DMPC/HistSuccDG/Chol 50:10:40,  
POPC/MoChol/DPPS/Chol 40:10:10:40,  
DPPC/DOTAP/DGSucc/Chol 20:10:30:40,  
DPPC/HistChol/Chol 50:10:40,

DPPC/HistSuccDG/Chol 40:20:40,  
DPPC/MoChol/DGSucc/Chol 20:10:30:40,  
POPC/HcChol/Chol 50:15:35,  
DPPC/HcChol/Chol 50:15:35,  
POPC/HistPS/Chol 50:15:35,  
DPPC/HistPS/Chol 50:15:35,  
POPC/AC/Chol 50:15:35,  
DPPC/AC/Chol 50:15:35,  
DPPC/HistChol/Chol 50:15:35,  
POPC/HistChol/Chol 50:15:35,  
DMPC/MoChol/DGSucc/Chol 20:10:30:40,  
POPC/HistSuccDG/Chol 50:15:35,  
DPPC/IsoHistSuccDG/Chol 50:15:35,  
DPPC/HistSuccDG/Chol 50:15:35,  
POPC/IsoHistSuccDG/Chol 50:15:35,  
DMPC/MoChol/DGSucc/Chol 20:10:30:40,  
POPC/MoChol/CHEMS/Chol 40:10:10:40,  
DMPC/HistChol/Chol 50:10:40,  
POPC/DOTAP/CHEMS/Chol 30:10:20:40,  
DMPC/HisChol/DGSucc/Chol 40:10:10:40,  
POPC/HisChol/CHEMS/Chol 40:10:10:40,  
DMPC/MoChol/CHEMS/Chol 40:10:10:40 ~~or and~~  
POPC/MoChol/DGSucc/Chol 30:20:10:40.

54. (Currently Amended) A method of treating a mammal with a drug comprising administering to the mammal the drug in the liposomal formulation[[s]] of claim 46.

55. (Currently Amended) The method of claim 954 wherein the mammal is a human.

56. (Currently Amended) The method of claim 954 wherein the liposomal formulation is administered parenterally for parenteral application, preferably intravenous application.

57. (Currently Amended) The method of claim 954, characterized in that it wherein the liposomal formulation includes one or more active substances.

58. (New) The liposomal formulation according to claim 46, wherein the proportion of cholesterol is 35 to 45 mole-% and the proportion of said mixtures is 15 to 45 mole-%.

59. (New) The liposomal formulation according to claim 46, wherein the oligonucleotide is a small interfering RNA.

60. (New) The liposomal formulation according to claim 46, wherein the oligonucleotide is a decoy oligonucleotide.